

AR
desferrioxamine (DFO); parathyroid hormone; anti-microbials, including, but not limited to anti-fungal agents; or any combination thereof.

IN THE CLAIMS:

Cancel claims 1-19 without prejudice.

Add claims 20-46 reading as follows:

20. A pharmacological composition comprising:

- (A) at least one biologically-active agent; and
- (B) at least one carrier compound having the formula



or a salt thereof



wherein Ar is a substituted phenyl or naphthyl;

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AB
R⁷ is selected from the group consisting of C₄ to C₂₀ alkyl, C₄ to C₂₀ alkenyl, phenyl, naphthyl, (C₁ to C₁₀)alkylphenyl, (C₁ to C₁₀)alkenylphenyl, (C₁ to C₁₀)alkylnaphthyl, (C₁ to C₁₀)alkenylnaphthyl, phenyl (C₁ to C₁₀)alkyl, phenyl (C₁ to C₁₀)alkenyl, naphthyl (C₁ to C₁₀)alkyl and naphthyl (C₁ to C₁₀)alkenyl;

R⁷ is optionally substituted with C₁ to C₄ alkyl, C₁ to C₄ alkenyl, C₁ to C₄ alkoxy, -OH, -SH and -CO₂R⁹ or any combination thereof;

R⁷ is optionally interrupted by oxygen, nitrogen, sulfur or any combination thereof;

R⁸ is selected from the group consisting of hydrogen, C₁ to C₄ alkyl, C₁ to C₄ alkenyl, hydroxy, and C₁ to C₄ alkoxy; and

~~R⁹ is hydrogen, C₁ to C₄ alkyl, or C₁ to C₄ alkenyl;~~
~~with the proviso that the compounds are not substituted with an amino group in the~~
~~position alpha to the acid group.~~

~~SAC
B1~~

21. The composition of claim 20, wherein Ar is substituted with at least one of
C₁-C₅ alkyl, C₂-C₄ alkenyl, -F, -Cl, -OH, -SO₂, -COOH or -SO₃H.

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~~A3~~
22. The composition of claim 21 wherein Ar is a substituted phenyl.

~~A3~~
3 23. The composition of claim 21, wherein Ar is a phenyl substituted with -Cl.

~~A3~~
4 24. The composition of claim 21, wherein Ar is a phenyl substituted with -F.

~~A3~~
5 25. The composition of claim 23, wherein R⁷ is selected from the group consisting
of C₄ to C₂₀ alkyl, (C₁-C₁₀ alkyl)phenyl, and phenyl (C₁ to C₁₀ alkyl).

~~A3~~
6 26. The composition of claim 23, wherein R⁷ is C₄-C₂₀ alkyl.

~~A3~~
7 27. The composition of claim 26, wherein R⁷ is not substituted or interrupted.

~~A3~~
8 28. The composition of claim 27, wherein R⁸ is hydrogen.

B2 29. The composition of claim 20, wherein the biologically active agent comprises at least one peptide, hormone, polysaccharide, mucopolysaccharide, carbohydrate, or lipid.

10 B2 30. The composition of claim 29, wherein the biologically active agent is a peptide.

11 B2 31. The composition of claim 29, wherein the biologically active agent is a mucopolysaccharide.

B3 SCB 32. The composition according to claim 20, wherein the biologically active agent comprises human growth hormone, bovine growth hormone, growth hormone-releasing hormone, an interferon, interleukin-1, interleukin-II, insulin, heparin, low molecular weight heparin, calcitonin, erythropoietin, atrial natriuretic factor, an antigen, a monoclonal antibody, somatostatin, adrenocorticotropin, gonadotropin releasing hormone, oxytocin, vasopressin, cromolyn sodium, vancomycin, desferrioxamine, parathyroid hormone, an antimicrobial, an antifungal agent or a combination thereof.

13 B3 33. The composition according to claim 32, wherein said biologically-active agent comprises human growth hormone, an interferon, insulin, heparin, low molecular weight heparin, calcitonin, erythropoietin, cromolyn sodium, parathyroid hormone, an antimicrobial or a combination thereof.

14 ~~34.~~ The composition according to claim 33, wherein said biologically-active agent comprises human growth hormone.

15 ~~35.~~ The composition according to claim 33, wherein said biologically-active agent comprises insulin.

AB *16* ~~36.~~ The composition according to claim 33, wherein said biologically-active agent comprises heparin.

17 ~~37.~~ The composition according to claim 33, wherein said biologically-active agent comprises low molecular weight heparin.

18 ~~38.~~ The composition according to claim 33, wherein said biologically-active agent comprises calcitonin.

19 ~~39.~~ The composition according to claim 33, wherein said biologically-active agent comprises cromolyn sodium.

20 ~~40.~~ The composition according to claim 33, wherein said biologically-active agent comprises parathyroid hormone.

D4 *526* ~~41.~~ A dosage unit form comprising

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Cont*

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(A) a pharmacological composition according to claim 20; and

(B) (i) an excipient,
(ii) a diluent
(iii) a disintegrant
(iv) a lubricant,
(v) a plasticizer
(vi) a colorant
(vii) a dosing vehicle, or
(viii) any combination thereof..

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42. A dosage unit form according to claim 41, comprising a tablet, a capsule, or a liquid.

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43. A dosage unit form according to claim 41, wherein said dosing vehicle is selected from the group consisting of water, 1,2-propane diol, ethanol, and any combination thereof.

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44. A method for preparing a pharmacological composition, said method comprising mixing:

(A) at least one biologically-active agent;
(B) at least one carrier compound having the formula



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wherein Ar is a substituted phenyl or naphthyl;

R5 Cost
from
R⁷ is selected from the group consisting of C₄ to C₂₀ alkyl, C₄ to C₂₀ alkenyl, phenyl, naphthyl, (C₁ to C₁₀)alkylphenyl, (C₁ to C₁₀)alkenylphenyl, (C₁ to C₁₀)alkyl)naphthyl, (C₁ to C₁₀)alkenyl)naphthyl, phenyl (C₁ to C₁₀)alkyl), phenyl (C₁ to C₁₀)alkenyl), naphthyl (C₁ to C₁₀)alkyl) and naphthyl (C₁ to C₁₀)alkenyl);

R⁷ is optionally substituted with C₁ to C₄ alkyl, C₁ to C₄ alkenyl, C₁ to C₄ alkoxy, -OH, -SH and -CO₂R⁹ or any combination thereof;

A B C D E F G H I J K L M N O P Q R S T U V W X Y Z
R⁷ is optionally interrupted by oxygen, nitrogen, sulfur or any combination thereof;

R⁸ is selected from the group consisting of hydrogen, C₁ to C₄ alkyl, C₁ to C₄ alkenyl, hydroxy, and C₁ to C₄ alkoxy; and

R⁹ is hydrogen, C₁ to C₄ alkyl, or C₁ to C₄ alkenyl;

with the proviso that the compounds are not substituted with an amino group in the position alpha to the acid group; and

(C) optionally a dosing vehicle.

45. A method for administering a biologically-active agent to an animal in need of said agent, said method comprising administering orally to said animal a composition as defined in claim 20.

46. A method for administering a biologically-active agent to a mammal in need of said agent, said method comprising administering orally to said mammal a composition as defined in claim 20.